

Microscopic scale conductivity as explanation of magnetotelluric results from the Alps of Western Switzerland

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SUMMARY

Recent MT soundings carried out in the Penninic Alps of Valais have shown the presence of a very good, outcropping conductor. Extremely high conductivity was attributed to the presence of graphite.

To verify this assumption, the electrical properties of borehole black shales were measured under simulated physical conditions (electrical frequency, hydrostatic confining pressure, internal fluid pressure, temperature). These measurements showed that under all physical conditions (electrical frequency, in the 0.005–200 Hz interval; hydrostatic confining pressure up to 39 MPa; internal fluid pressure up to 23 MPa; temperature up to 180 °C) one of the samples studied was very conductive (resistivity less than 2 Ω m under all physical conditions). Interestingly, despite similar macroscopic aspect, other samples from a nearby borehole were found to be only slightly conductive. Chemical, mineralogical and petrographic investigations revealed that the enhanced electrical conductivity is mostly due to textural characteristics (such as grain size and carbon film distribution at the grain boundaries) rather than to chemical differences.

Key words: Alps, electrical conductivity, magnetotellurics, graphite

1 INTRODUCTION

Magnetotelluric soundings performed in various areas have indicated the presence of low-resistivity structures in the upper and middle crust. Among these areas there are the following European sites, the Lapland granulite belt (Korja *et al.* 1996), the Carpathians (Praus *et al.* 1990), Scotland and Ireland: Precambrian collision belt (Whelan *et al.* 1990), the Pannonian basin (Adam & Bielik 1998) the EGT Central segment (ERCEUGT-Group 1992).

The cause for enhanced conductivity is often thought to be related to the presence of carbon in the form of graphite. However, the nature of the good conductor often remains an open issue in the absence of direct investigation techniques, although extremely high conductivity values may be explained only by the presence of graphite and meta-anthracite-bearing rocks (Jödicke 1992). In general, no simple cause is found, and candidates such as saline fluids saturating the pores, or inter-connected graphite are often invoked (Frost *et al.* 1989; Glover & Vine 1992; Jones 1992; Glover & Vine 1994; Glover & Vine 1995; Shankland *et al.* 1997; Roberts *et al.* 1999).

Schnegg (1998) carried out a magnetotelluric survey in the penninic part of the Swiss Alps, canton Valais, western Switzerland. Multidimensional modelling of the resulting data

(Schnegg 1993, 1999) revealed the existence of a very conductive slab embedded in a much more resistive rock matrix. The shape of the slab correlates well with reflecting horizons from an existing deep seismic reflection profile acquired across the Western Swiss Alps, that dips 35° SSE (Pfiffner *et al.* 1997). Furthermore, the modelling required that the northernmost edge of the slab was close to or at the surface. This edge was effectively found in the form of black shales.

VLF profiles have been measured in the area of the outcrop. Low resistivity cover (<5 Ω m, challenging values for the VLF receiver) was found in this area. The occurrence of the outcrop provided an opportunity to study the cause of enhanced electrical conductivity in the upper- and middle crust using lab investigations of samples from two boreholes located in the area where MT data modelling indicated proximity to the body responsible for the high conductivity.

2 GEOLOGICAL SETTING

The region surveyed by MT is located in canton Valais and covers an area of 3200 km² (Fig. 1). The Rhone Valley forms the contact between two domains: the Helvetic Zone to the NNW, and the Penninic domain to the SSE; all the MT field work was done on the latter. These domains overlay a pre-Triassic

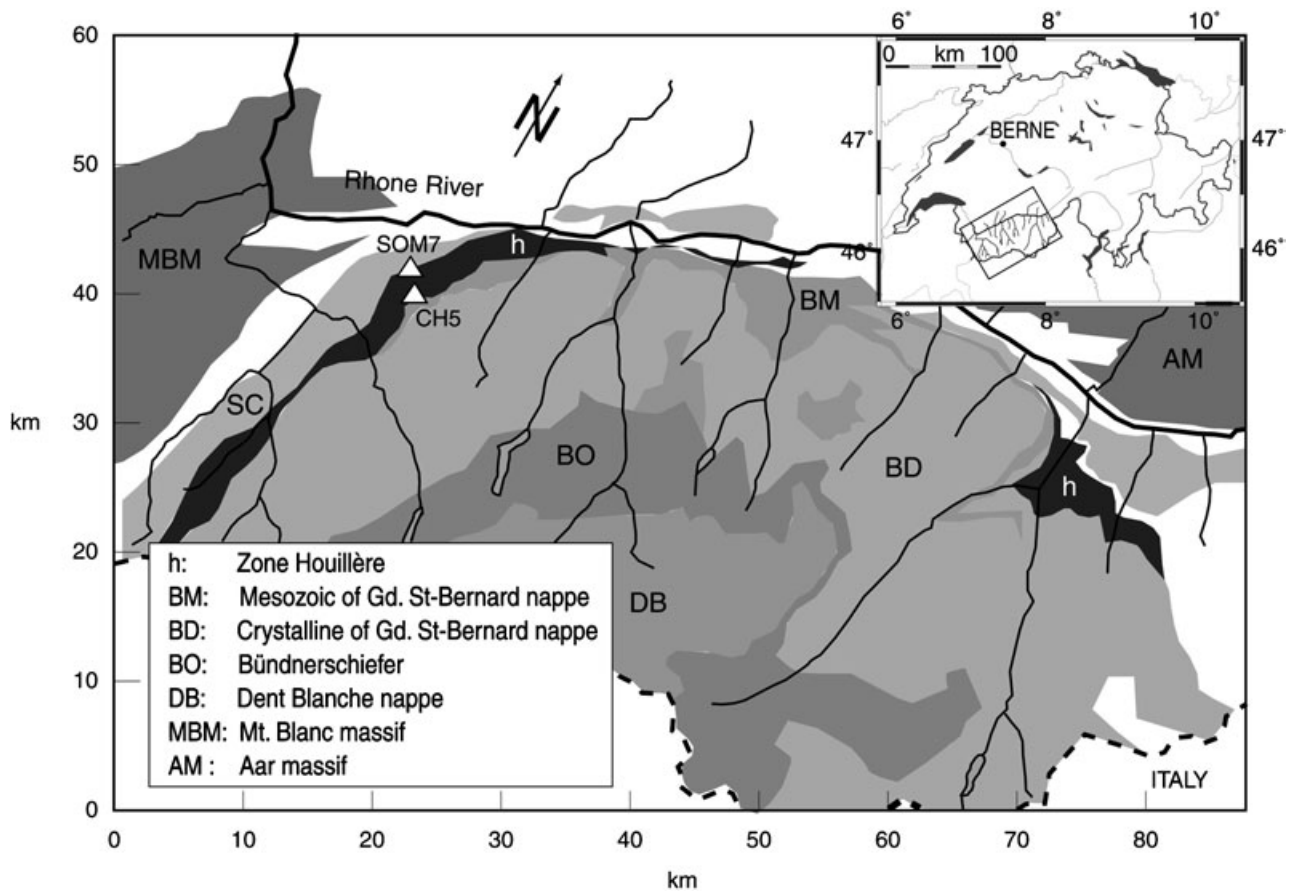


Figure 1. Tectonic map of the studied area showing the location of the boreholes SOM7 and CH5 (triangles).

basement complex, which has been modified by the Variscan orogeny. Mid-Cretaceous to Pliocene compressional movements acting on the Penninic ocean resulted in the development of the Penninic Alps. As many as four tectonic units are stacked over one another moving NNW to SSE. These are (from bottom to top) (Escher *et al.* 1997): (1) a thick sedimentary series (2) the Grand St-Bernard nappe, with its northernmost constituents, the Upper Carboniferous of the Zone Houillère and the Permo-Carboniferous of the Stalden unit (1–2 km of dark schists and anthracite-bearing slates) representing potentially good conducting layers due to possible graphite content (3) the Schiste lustré Mesozoic sediments and (4) the crystalline basement rocks of Dent Blanche nappe. To the SW and NE, two crystalline, pre-Permian autochthonous blocks limit the area of study (Fig. 1).

Cylindrical core samples (diameter: 78.5 mm) studied as part of this investigation were obtained from two boreholes drilled in the Zone Houillère (see location on Fig. 1). This segment forms the lower, external part of the Grand St-Bernard nappe. The samples consisted of sandstone and hand-staining black shales. The Zone extends laterally over more than 200 km and

represents the detrital filling of a Permo-Carboniferous graben system which was detached from its basement by alpine movements. The location of CH5 and SOM7 boreholes is given in Table 1.

3 EXPERIMENTAL FIELD METHODS: MAGNETOTELLURIC SURVEY, SEISMIC PROFILE AND MODEL

MT and GDS measurements were taken over an area of 3200 km² limited to the north by the Rhone river and to the South by the Swiss–Italian border. The location of the sites, the magnetotelluric data and the modelling results can be found elsewhere (Schnegg 1998, 1999). 2- and 3D modelling revealed the existence of a very low resistive (0.2 Ω m), 1 km-thick slab embedded in a resistive, featureless 1300–3000 Ω m rock matrix. The slab plunges in the SSE direction below the Alps with an average dip of 35° and follows the shape of the Alpine arc. With a total length of 20 km, its lower tip reaches a depth of 8 km. At its NNW edge, it crops out at the surface in the middle of the Houillère Zone.

Table 1. Location of the boreholes.

Borehole	Abbr.	Coordinates	Elevation (m)	Depth (m)
Les Etablons	CH5	7°14'6"E/46°7'44"N	2048	318
Som de Pro window	SOM7	7°13'13"E/46°8'39"N	1457	31.5

Comparison with existing seismic profiles (Pfiffner *et al.* 1997) shows very good agreement between the slope of the conducting feature in the MT model and the seismic reflectors. The plunging, very conducting layer can be interpreted as the southward continuation of the Zone Houillère outcrop.

4 LABORATORY EXPERIMENTAL METHODS

To constrain the interpretation of the MT and GDS results, we studied the electrical properties of 3 representative samples from two boreholes located 4–5 km to the N of Verbier (SOM7 and CH5, see Fig. 1), on the NE flank of the Pierre Avoi peak. Both boreholes were drilled into the Houillère Zone. The samples were examined using chemical, mineralogical and petrographical methods.

Three 36 mm diameter high schistosity black shale samples were investigated; two samples from borehole CH5 (CH5 \perp : schistosity perpendicular to cylinder axis, length: 40 mm; CH5//: schistosity parallel to cylinder axis, length: 70 mm) and 1 sample from borehole SOM7 (schistosity 45° with respect to cylinder axis, length: 40 mm).

4.1 Experimental electrical resistivity methods

The electrical properties of the samples were studied under simulated physical conditions (temperature T up to 180 °C; confining hydrostatic pressure P_e up to 39 MPa; internal pore fluid pressure P_i up to 23 MPa; current frequency ranging between 0.005 and 200 Hz). This study was performed using the experimental method described by Losito *et al.* (1991). The samples were studied without drying before the experimental cycles, because their cleavage and mineralogical composition make them very fragile. The 2-electrode array technique was used for the smallest samples (CH5 \perp and SOM7), whereas the standard 4-electrode array technique was used for the largest sample (CH5//). As is well known the 2-electrode technique cannot keep the electrode polarization apart from the sample polarizations. The use of a 4-electrode array for the CH5 \perp sample made evaluation of the electrode polarization of this sample possible and allowed us to extrapolate this information to other samples. The electrodes at the cylinder bases consisted of stainless steel plates (36 mm diameter, fully covering the cylinder bases) and are used as current electrodes for the 4-electrode array, but as current and voltage electrode for the of 2-electrode array. Good electrode to sample contact is achieved with a porous surface. The electrodes connect to stainless steel capillary tubes used to inject and pressurize the sample fluids, allowing homogeneous diffusion of the fluid. The voltage electrodes of 4-electrode array consisted of 0.3 mm stainless steel wire. They were installed in an *ad hoc* sample canal far away from the current electrodes. The electrodes have been accurately tested in the whole frequency interval used for the experiments under dry and hydrated conditions. Polarization phenomena were completely absent. The samples and their electrodes were enclosed in a silicone polymer, selected according its elastic properties under the physical experimental conditions (max. working temperature: 240 °C).

The electrical parameters were obtained using a pure sinusoidal waveform as the input signal; the frequency interval has been selected to correspond with the MT frequencies,

because various polarization phenomena may occur at every frequency interval. To describe the electrical behaviour of the rock, the real part of the complex resistivity ρ (Ωm), the permittivity ϵ (F/m) and the loss tangent $\tan\delta$ were automatically computed (Losito 1989) for each selected frequency. The total harmonic distortion rate (THD per cent) of the output signals was used to filter out the noise and the non-linear component of the output signals (Losito 1989). The complex resistivity parameters were calculated by using only the first harmonics of the input and output signals. To simulate the underground physical conditions, the thermo-baric cycles have been carried out following a rigorous procedure. Under these operating conditions, the heating produces permanent petro-physical alterations, mainly related to water and OH^- content and distribution (Losito *et al.* 1991). Pressure can also produce permanent mechanical alterations (micro and macro cracks). We assume that every change (mineralogical or structural) is non-reversible, but the effect of the temperature on the conductivity is more important during the experiment interval. Consequently, we decided to submit the samples first to pressure changes, followed by temperature variations. Before launching the thermo-baric cycle, the sample was hydrated inside the vessel by injection of pure water through a capillary tube–porous electrode system. At the same time the hydration status was monitored by observing (1) resistivity changes of the sample and (2) water output through a second capillary tube–porous electrode system. In the following section, the experimental results will be analytically described, by comparison of the electrical behaviour of the samples with respect to variations of a main conditioning parameter (frequency, external confining pressure P_e , or internal fluid pressure P_i) at $T = 20, 50, 80, 110, 150$ and 180 °C.

4.1.1 Time dependence

The samples, at natural water content, sealed by the silicon glue, were hydrated inside the vessel with deionized water using the porous electrode–capillary tube system. The resistivity at frequency 0.1 Hz has been measured during the initial flow of the injected fluid P_i , controlling the confining pressure P_e at the same time (SOM7 sample: $P_i = 3$ MPa, $P_e = 6$ MPa; CH5 samples: $P_i = 5$ MPa, $P_e = 9$ MPa). Due to the very low hydraulic conductivity of the samples the applied pressure of the pore fluid did not allow the full saturation of the sample (no fluid observed outside the unconnected capillary tube after 3000 min). The samples were dried after the thermo-baric cycle. The water loss resulted in less than 0.1 per cent weight change, which is consistent with measurements of the mass of water adsorbed.

The laboratory measurements illustrate the high resistivity contrast between the two CH5 samples and the SOM7 sample (see. Table 2). In fact the CH5 samples appeared to be 3 orders of magnitude more resistive than the SOM7 sample, in spite of

Table 2. Effects of hydration cycle on sample resistivities.

Sample	ρ (Ωm) ($t = 0.0$ min)	ρ (Ωm) ($t = 3000$ min)
CH5//	2280.0	744.0
CH5 \perp	7900.0	6040.0
SOM7	2.3	1.39

a very similar macroscopic appearance. These initial measurements also indicate a strong anisotropy effect (Lastovickova *et al.* 1993) in the CH5 samples. The variation of resistivity with time is slow and small. The observed continuous resistivity decrease must be attributed to the small quantity of water injected, which obviously acts more in the sample CH5//, where the layers are parallel to both water and current flux and thus, allow better formation of surface conduction paths.

4.1.2 Frequency dependence

The frequency dependence was measured in the 0.005–200 Hz range, under the internal pore fluid pressures, hydrostatic pressures, and temperatures as indicated in Figs 2(a), 3(a), and 4(a).

In general the resistivity $\rho(\omega)$ shows small linear dispersive phenomena, more pronounced at lower frequencies. In particular

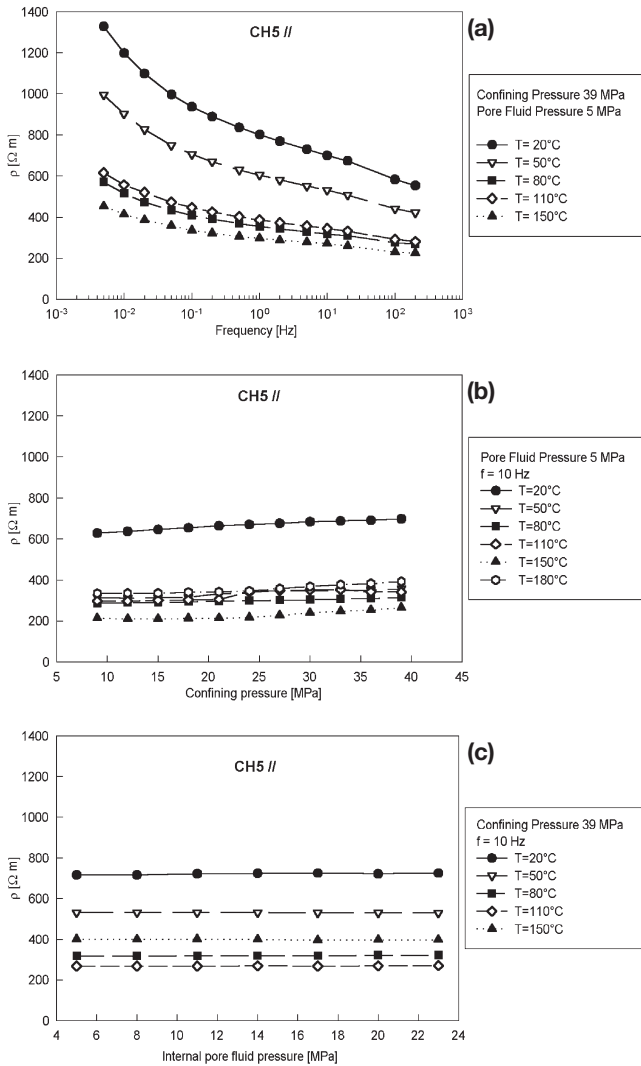


Figure 2. Resistivity ρ (Ωm) of CH5//sample (CH5 borehole, 318 m depth, layers parallel to the cylinder axis) at the given temperatures, under different simulated physical conditions; (a) as function of frequency; (b) as function of hydrostatic confining pressure; (c) as function of internal pore fluid pressure.

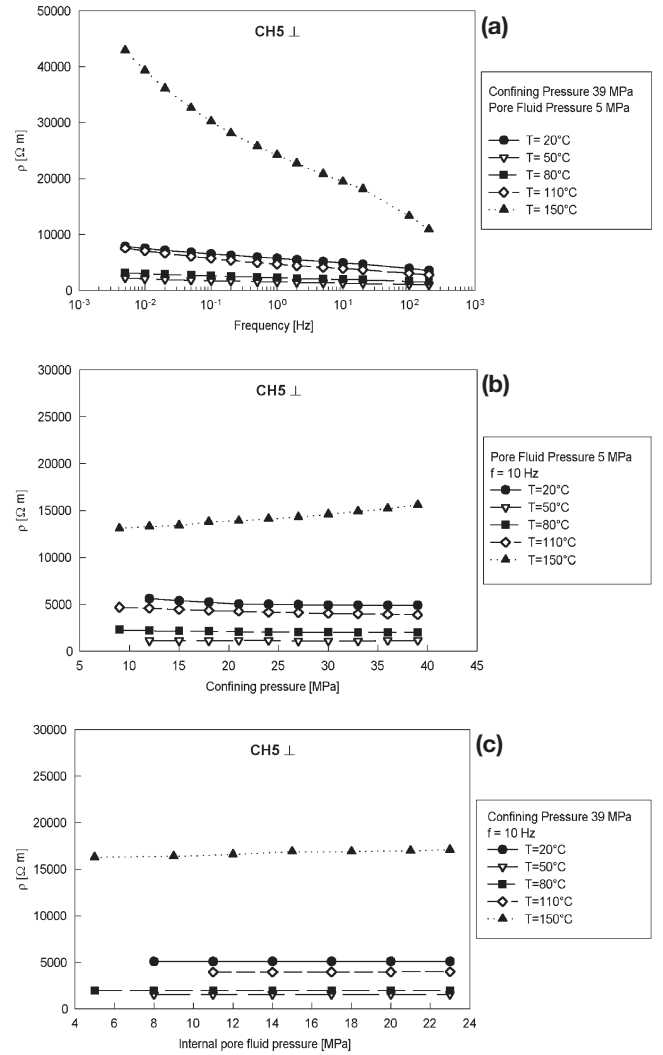


Figure 3. Resistivity ρ (Ωm) of CH5⊥ sample (CH5 borehole, 318 m depth, layers perpendicular to the cylinder axis) at the given temperatures, under different simulated physical conditions: (a) as function of frequency; (b) as function of hydrostatic confining pressure; (c) as function of internal pore fluid pressure.

for the CH5 samples a noticeable frequency dispersion of ρ at very low frequencies (0.005–5 Hz, Figs 2a and 3a) is apparent; this dispersion must be attributed to phyllosilicate minerals (Losito & Muschietti 1987). A second polarization, confirmed by $\tan \delta(\omega)$ measurements at all temperatures and pressures, is present at frequencies $f > 0.1$ Hz and can be attributed to a metallic phase, such as pyrite, or possibly due to graphite (unpublished test on pure dry graphite shows frequency dispersion in the 20–200 Hz range). The SOM7 sample (Fig. 4a) shows frequency dispersion of ρ (of about 20 per cent) to occur in the 0.1–100 Hz interval at $T = 20^\circ\text{C}$ (at this temperature the confined pressure was 6 MPa). This frequency dispersion is similar to the high frequency dispersion of CH5 samples, but more marked. The frequency trend of SOM7 ρ at $T = 20^\circ\text{C}$ shows a jump at $f = 0.1$ Hz, that could be attributed to a small observed increase of the internal pore fluid pressure (1 MPa). At higher temperatures we have minor frequency dispersion of ρ respect to $T = 20^\circ\text{C}$ curve in a higher frequency range (10–100 Hz). This small frequency dispersion is confirmed also

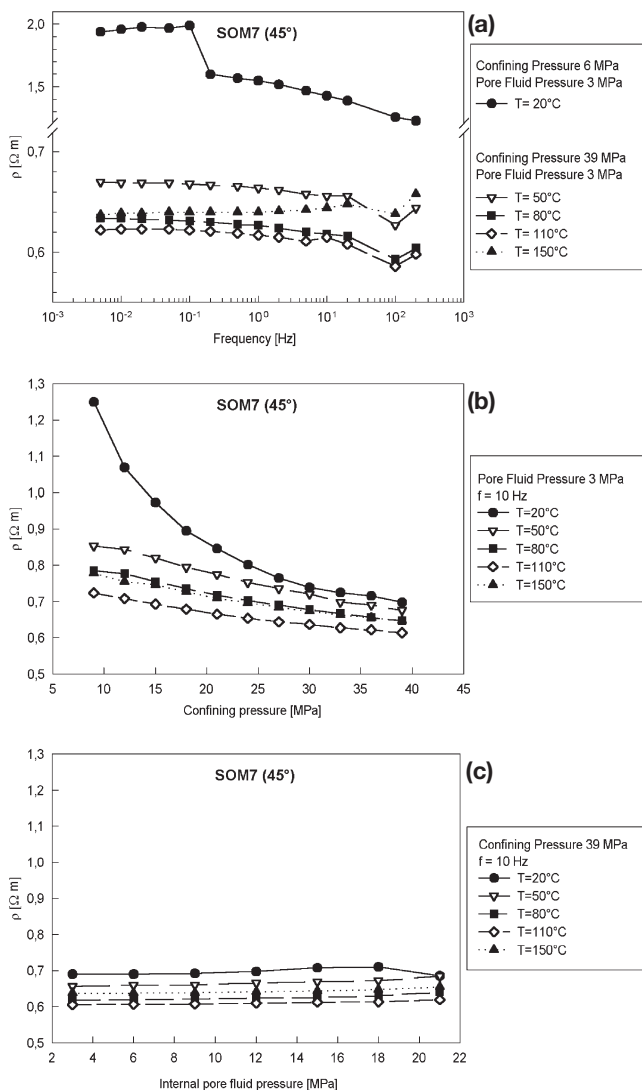


Figure 4. Resistivity ρ (Ωm) of SOM7 sample (SOM7 borehole, 31.5 m depth, layers 45° with respect to the cylinder axis) at the given temperatures, under different simulated physical conditions: (a) as function of frequency; (b) as function of hydrostatic confining pressure; (c) as function of internal pore fluid pressure.

by $\tan \delta(\omega)$, that is 0.35 at 20 Hz. The ρ behaviour at higher temperatures can be attributed to linear dispersive phenomena, such as those induced by graphite.

4.1.3 Temperature dependence

In general resistivity decreases with temperature up to 80°C . This behaviour depends on the electrokinetic activity of the ions inside the fluid. From Fig. 4, it is evident that the resistivity decrease with temperature of the SOM7 sample is not expected. We attribute this to a higher carbon content compared to the CH5 samples. The curves of SOM7 for $T=20^\circ\text{C}$ are quite different from others; the cause could be the low hydraulic permeability of this sample, which cannot produce high electrokinetic effects at higher temperatures. At temperatures above 100°C the vapour effect is dominant and the resistivity depends on the joint effects of pressures (confining and pore fluid), temperature and sample foliation with respect to the probe axis,

as shown with the CH5 samples (Figs 2 and 3). The resistivity increase at high temperature may also depend on the OH^- layers inside the phyllosilicates, as their movement can be activated by thermal energy.

It is important to note that the whole experimental cycle has been applied to each sample, so the resistivity decrease can be interpreted as an effect of repeated pressure cycling at increasing temperatures and should only be understood as relative conductivity measurements with respect to the initial values. However we prefer to show non-normalized resistivities.

4.1.4 External Confining Pressure dependence

The confining pressure produces opposite effects on electrical behaviour (measured at 10 Hz) of CH5 and SOM7 samples: resistivity shows small increases in the CH5//sample and in the CH5 \perp only at $T=150^\circ\text{C}$, but resistivity decreases in sample SOM7; this behaviour can be attributed both to different grain sizes and carbon content. More specifically, for CH5 samples (Figs 2b and 3b) the foliation affects the electrical resistivity (with an anisotropy factor of more than 10). The increase of external confining pressure produces increase of the resistivity in sample CH5 \perp (layers perpendicular to fluid injection) only at high temperatures (150°C). The resistivity of sample CH5// increases slightly as function of confining pressure at all temperatures, or is constant. This is related to the vapour action upon layering (microcracks). For the SOM7 sample (Fig. 4b), the resistivity decrease with increasing confining pressure is apparent at all studied temperatures; this behaviour can be attributed to the reconnection of conductive minerals, such as graphite. The dispersion of ρ at high frequencies under all temperatures appears to confirm this hypothesis.

4.1.5 Pore Fluid Pressure dependence

For all samples, increasing the pore fluid pressure had no relevant effect on the resistivity. The main cause of this behaviour is the very low hydraulic conductivity of the black shales, as indicated by the hydration cycles (Table 2). Consequently the pressurized pore fluid does not have sufficient access to the rock specimen to enter into connected pores and the pore fluid pressure cannot reach the equilibrium before the measurements were taken. It is possible that experiments of many months could solve the problem, using the working pressures described.

4.1.6 Concluding remarks

The results of the laboratory measurements described require further explanation of the cause of the resistivity difference between the samples from the two boreholes, measured under identical physical conditions. The laboratory results confirm the MT data, but do not allow an immediate petrophysical interpretation based on macroscopic petrographical evidences. Principal selected candidates in solving the problem were the carbon content and its connectivity. Nevertheless it is also important to know the phyllosilicate content and identify the metallic minerals (Losito & Muschietti 1998).

To investigate this issue further, chemical and petrographical investigations were carried out. These are described in the following section.

5 PETROGRAPHIC AND MINERALOGICAL STUDIES

The SOM7 and CH5 samples are black shales with similar appearance in hand specimen (no clear differences in texture, grain size and colour). Petrographic, chemical and mineralogical investigations were carried out to understand their different electrical behaviour. Attention was mostly paid to the C-content, the occurrence of C-minerals and to the rock texture (distribution of the C film).

The petrographic studies were carried out starting from four petrographic sections for each sample; the sections have been selected in different orientations with respect to the main foliation (one parallel, one perpendicular and two oblique). The SOM7 and CH5 samples appear to be mostly made of quartz and finely intergrown phyllosilicates. By comparing the corresponding petrographic sections of SOM7 and CH5 (i.e. sections in the same orientation with respect to the main foliation), only slight textural difference is observed; in particular, foliation is wider and broader in CH5 than in SOM7; moreover, grain size of the SOM7 sample appears to be smaller than in CH5. This evidence has been recently confirmed by crystallite size analyses, performed for samples SOM7 and CH5 at the Department of Earth Sciences in Florence. Reasonably, grain size could influence the C-film connectivity (in particular, the lower the grain size, the higher the connectivity).

The whole-rock chemical composition of SOM7 and CH5 have been obtained by X-ray fluorescence (XRF) spectrometry (Philips PW1414). Samples have very similar compositions (Table 3): main differences are in Na_2O , MgO , K_2O , Ba and Sr.

These data have been integrated by recent elemental measures at the Chemical Laboratory of Florence University, revealing that the C-content is 3.3 per cent for SOM7 and 1.34 per cent for CH5. The observed slight difference in the C-content cannot explain the difference in resistivity. The absence of a direct correlation between rock C-content and conductivity has been already observed by Korja *et al.* (1996).

X-ray diffraction (XRPD) patterns on whole-rock powders have been collected on a Philips PW1710 diffractometer, working

at 45 kV and 25 mA (radiation: $\text{CuK}\alpha$): XRPD patterns provide a list of characteristic d_{hkl} spacing, from which it is possible to determine the main rock-forming minerals. The experimental patterns of both SOM7 and CH5 revealed the presence of quartz, micas and chlorite (except for four very weak peaks, which cannot be related to any of the above minerals). Graphite was not detected, possibly due to its low crystallinity (producing very weak and broad diffraction) and to the interference between graphite peaks and those of the other present minerals.

The petrographic sections of SOM7 and CH5 have been examined at the scanning electron microscope (SEM, Philips XL30 working at 20 kV). An X-ray energy dispersive spectrometer (EDS–EDAX DX4) has been used for mineral chemistry determinations. Since the X-ray detector is equipped with an ultra-thin window, it was possible to detect also light elements such as carbon. Moreover, owing to their high conductivity, neither SOM7 nor CH5 samples needed to be carbon-coated, as usual in electron microscopy: consequently, the presence of carbon minerals has been unequivocally evidenced from the corresponding EDS spectra.

In order to identify graphite in SEM images, mostly Back Scattered Electrons (BSE) have been used for image formation. In fact, contrast in BSE images is related to the mean atomic number; minerals with a low mean atomic number are dark, whereas minerals with higher mean atomic number are more or less bright. In SOM7 and CH5, dark areas correspond to graphite. Coarse grained grey crystals are mostly quartz (rarely feldspar), grey fine grained aggregates correspond to phyllosilicate intergrowths, white radial aggregates are formed by chloritoid, white euhedral crystals are apatite and opaque minerals (Figs 5 and 6). The BSE images show very well the differences in texture and in graphite distribution between SOM7 and CH5. In sample CH5, two different foliations are often evident (Fig. 5). Graphitic aggregates typically occur in thin layers (up to 50–100 μm wide) parallel to the foliations. These layers are irregularly spaced within the sample, separating wide areas mainly formed by silicates. Graphite is not homogeneously distributed along the layers, but appears as a dark continuum on the optical microscope. The size of the graphite aggregates, always elongated parallel to the foliation, is variable. When two foliations are evidenced, coarse grained aggregates (up to 200 μm) are associated with the first foliation, whereas finer aggregates occur parallel to the second foliation.

In SOM7 graphite layers are finer and homogeneously distributed in the sample; graphite surrounds the silicate crystals, thus producing a continuous film (Fig. 6). We suppose that the high conductivity of SOM7 is due to the finely dissembling distribution of graphite.

The transmission electron microscopy (TEM) provides extremely wide information; in particular we can simultaneously obtain enlarged images of the sample, the corresponding diffraction pattern and the chemical composition, with a spatial and analytical resolution higher than with other conventional instruments. The availability of both chemical and diffraction data from very small C-rich areas allows the rigorous identification of graphite. A TEM microscope (Philips 400T, 120 kV), with a nominal point to point resolution of 4 Å was used to carry out the present study. Samples have been prepared by the ion-thinning method, using a Gatan Dual Ion Mill 600. Electron diffraction patterns are referred to areas as small as *c.* 0.5 μm in diameter (SAED = selected area electron diffraction). An EDAX DX4 energy dispersive spectrometer (EDS), equipped

Table 3. Chemical composition of studied samples. * ($\text{FeO} + \text{Fe}_2\text{O}_3$).

	SOM7	CH5	wt% ppm	SOM7	CH5
H_2O	5.75	4.98	Nb	21	11
Na_2O	2.73	1.53	Zr	203	275
MgO	0.70	1.33	Y	34	43
Al_2O_3	23.12	22.03	Sr	334	84
SiO_2	58.82	58.60	Rb	126	250
P_2O_5	0.09	0.05	Ni	18	29
K_2O	2.44	5.22	Cr	54	86
CaO	0.17	0.12	V	93	134
TiO_2	0.76	1.02	Ba	434	1173
MnO	0.05	0.04	La	62	53
Fe_2O_3^*	5.37	5.07	Ce	110	91
			Pb	31	44
			Co	17	17
			Th	30	30
			U	8	14
			Zn	83	90

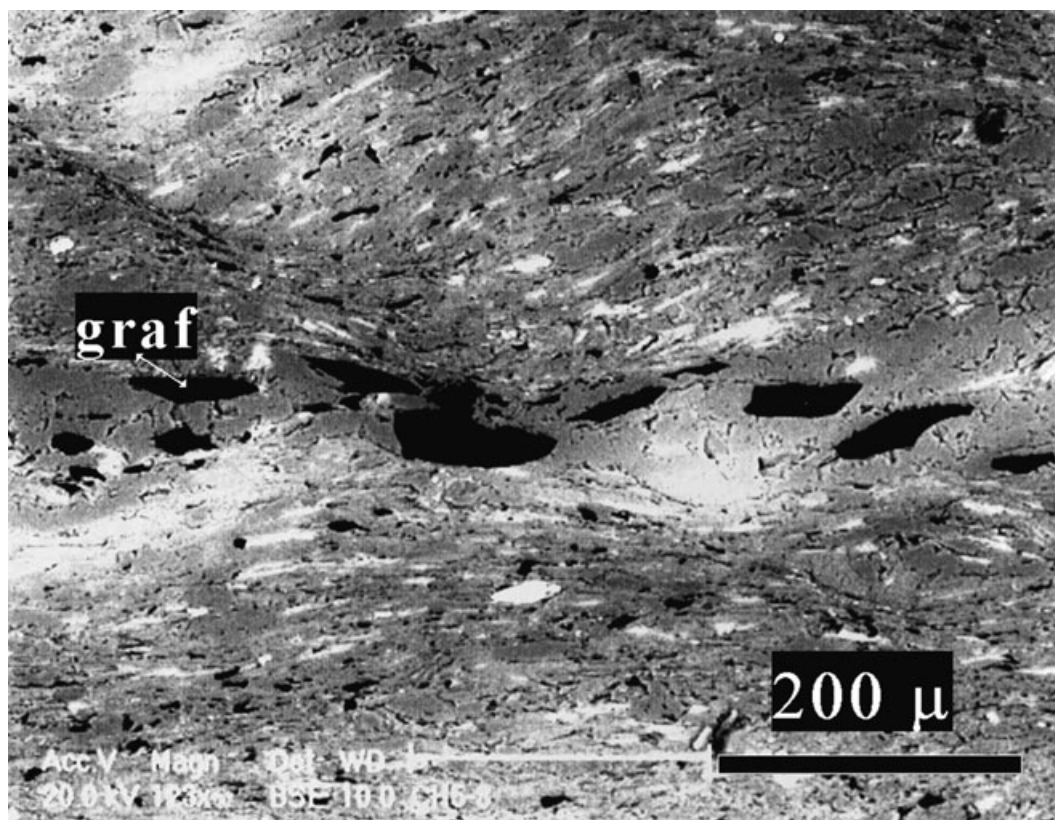


Figure 5. SEM image of a sample from borehole CH5. Section perpendicular to the schistosity. Graphitic areas are denoted by the mention 'graf'.

with ultra thin window, also provides chemical data for carbon. Chemical data have been calculated according to the method of Cliff & Lorimer (1975).

Graphite has been clearly identified by TEM; it forms irregular microcrystalline aggregates, whose maximum dimensions are approximately $5 \times 2 \mu\text{m}$. EDS analyses revealed that only carbon is present in these aggregates. Graphitic aggregates are round-shaped and slightly elongated according to the main foliation. Graphite single crystals are featureless and not clearly evident in images: anyway, crystal size appears to be below 500 \AA . Electron diffraction patterns of aggregates are ring-shaped (thus indicating a polycrystalline structure), weak and

diffuse. Only a few diffraction rings are present, but the corresponding d_{hkl} can be attributed to graphite. At the TEM scale, there are no clear differences between SOM7 and CH7 graphitic aggregates. Moreover, we can argue that the low crystallinity of graphite does not strongly influence the sample conductivity.

CONCLUSIONS

In a former work, magnetotelluric (MT) and geomagnetic depth soundings (GDS) have revealed the existence of a very high conducting layer below the Penninic Alps of Valais. Outstanding geometrical fit of the conductivity model with seismic lines helped identify the lithology responsible for the enhanced conductivity (Zone Houillère). This layer crops out at the surface, making it possible to check in the laboratory the reality of this electric conductor. The electrical properties of black shales from the boreholes, measured under simulated physical conditions showed that under all physical conditions the sample from one borehole (SOM7) was very conductive ($\rho < 2.0 \Omega\text{m}$), despite a close visual resemblance to the much less conductive samples from the second borehole nearby (CH5, $\rho > 300 \Omega\text{m}$). Moreover, the laboratory results (SOM7) confirmed the field data, but could not suggest an immediate petrophysical interpretation.

The principal explanations for the difference were differences in carbon content and variations in conductive mineral connectivity. It was also important to know the phyllosilicate content and to identify the metallic minerals. To these ends chemical and petrographical investigations were performed, using

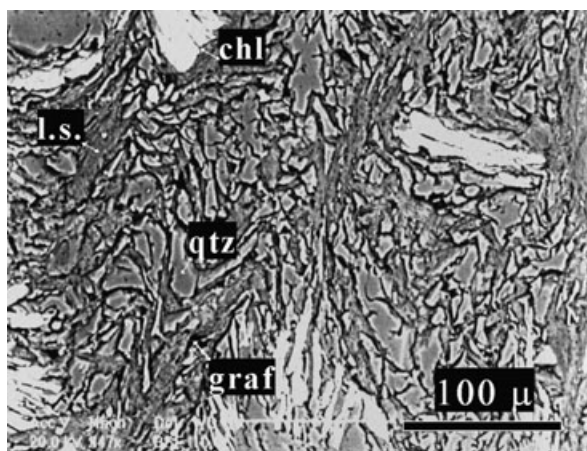


Figure 6. SEM image of a SOM7 sample. Section parallel to the schistosity. Graphitic areas are denoted by the mention 'graf'.

both classical (optical microscope) and sophisticated (TEM, SEM). These investigations revealed that the cause of the electrical conductivity was not of chemical origin, but petrographical and related to structure layering, the grain size of phyllosilicates, and the presence of a carbon film at the grain boundaries. Grain size and layering are typical of fluvial deposits, leading to stratigraphic heterogeneities such as lenses and channels. Bed continuity is not insured over large distances. Under these circumstances, it is not surprising to find large conductivity variations between neighbouring wells, and even within the same rock core.

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